

EXPRESS MAIL LABEL NO.:

EVO64374867US

DATE: March 25, 2002

JC02 Rec'd PCT/PTO 25 MAR 2002


FORM PTO 1300 (REV 11 2000)		U.S. DEPARTMENT OF COMMERCE, PATENT AND TRADEMARK OFFICE	ATTORNEY'S DOCKET NUMBER OYJALO-011
TRANSMITTAL LETTER TO THE UNITED STATES DESIGNATED/ELECTED OFFICE (DO/EO/US) CONCERNING A FILING UNDER 35 U.S.C. 371		U.S. APPLICATION NO. (if known, see 37 CFR 1.5) 10/089022	
INTERNATIONAL APPLICATION NO. PCT/FI00/00855	INTERNATIONAL FILING DATE 4 October 2000	PRIORITY DATE CLAIMED 4 October 1999	
TITLE OF INVENTION PROCESS FOR PREVENTING SCALING OF HEAT TRANSFER SURFACES			
APPLICANT(S) FOR DO/EO/US Asko Paakki, Päivi Uusitalo, and Mikael Svedman			

Applicant herewith submits to the United States Designated/Elected Office (DO/EO/US) the following items and other information.

- ☒ This is a **FIRST** submission of items concerning a filing under 35 U.S.C. 371.
- ☐ This is a **SECOND** or **SUBSEQUENT** submission of items concerning a filing 35 U.S.C. 371
- ☒ This is an express request to begin national examination procedures (35 U.S.C. 371 (f)). The submission must include items (5), (6), (9) and (21) indicated below.
- ☐ The US has been elected by the expiration of 19 months from the priority date (PCT Article 31).
- ☒ A copy of the International Application as filed (35 U.S.C. 371 (c)(2))
 - ☐ is attached hereto (required only if not communicated by the International Bureau).
 - ☒ has been communicated by the International Bureau.
 - ☐ is not required, as the application was filed in the United States Receiving Office (RO/US).
- ☐ An English language translation of the International Application as filed (35 U.S.C. 371 (c)(2)).
 - ☐ is attached hereto.
 - ☐ has been previously submitted under 35 U.S.C. 154(d)(4).
- ☒ Amendments to the claims of the International Application under PCT Article 19 (35 U.S.C. 371 (c)(3))
 - ☐ are attached hereto (required only if not communicated by the International Bureau).
 - ☐ have been communicated by the International Bureau
 - ☐ have not been made; however, the time limit for making such amendments has NOT expired.
 - ☒ have not been made and will not be made.
- ☐ An English language translation of the amendments to the claims under PCT Article 19 (35 U.S.C. 371 (c)(3)).
- ☒ An oath or declaration of the inventor(s) (35 U.S.C. 371 (c)(4)). (Executed)
- ☐ An English language translation of the annexes to the International Preliminary Examination Report under PCT Article 36 (35 U.S.C. 371 (c)(5))

Items 11 to 20 below concern document(s) or information included:

- ☐ An Information Disclosure Statement under 37 CFR 1.97 and 1.98.
- ☒ An assignment document for recording. A separate cover sheet in compliance with 37 CFR 3.28 and 3.31 is included.
- ☐ A **FIRST** preliminary amendment.
- ☐ A **SECOND** or **SUBSEQUENT** preliminary amendment.
- ☐ A substitute specification
- ☐ A change of power of attorney and/or address letter
- ☐ A computer-readable form of the sequence listing in accordance with PCT Rule 13ter.2 and 35 U.S.C. 1.821 - 1.825.
- ☒ A second copy of the published international application under 35 U.S.C. 154(d)(4).
- ☐ A second copy of the English language translation of the international application under 35 U.S.C. 154(d)(4).
- ☒ Other items or information Copy of International Preliminary Examination Report, Copy of International Search Report, One (1) Sheet Formal Drawings

U.S. APPLICATION NO. (if known, see 37 CFR 1.5) <div style="font-size: 24pt; font-weight: bold; text-align: center;">10/089022</div>	INTERNATIONAL APPLICATION NO. PCT/FI00/00855	ATTORNEY'S DOCKET NUMBER OYJALO-011	
21. <input checked="" type="checkbox"/> The following fees are submitted:		CALCULATIONS PTO USE ONLY	
BASIC NATIONAL FEE (37 CFR 1.492 (a) (1) - (5)): <input checked="" type="checkbox"/> Neither international preliminary examination fee (37 CFR 1.482) nor international search fee (37 CFR 1.445(a)(2)) paid to USPTO and International Search Report not prepared by the EPO or JPO \$1040.00 <input type="checkbox"/> International preliminary examination fee (37 CFR 1.482) not paid to USPTO but International Search Report prepared by the EPO or JPO \$890.00 <input type="checkbox"/> International preliminary examination fee (37 CFR 1.482) not paid to USPTO but international search fee (37 CFR 1.445(a)(2)) paid to USPTO \$740.00 <input type="checkbox"/> International preliminary examination fee (37 CFR 1.482) paid to USPTO but all claims did not satisfy provisions of PCT Article 33(1)-(4) \$710.00 <input type="checkbox"/> International preliminary examination fee (37 CFR 1.482) paid to USPTO and all claims satisfied provisions of PCT Article 33(1)-(4) \$100.00			
ENTER APPROPRIATE BASIC FEE AMOUNT = Surcharge of \$ _____ for furnishing the oath or declaration later than <input type="checkbox"/> 20 <input type="checkbox"/> 30 months from the earliest claimed priority date (37 CFR 1.492 (c)).		\$ 1,040.00 \$ _____	
CLAIMS	NUMBER FILED	NUMBER EXTRA	RATE
Total claims	18-20 =		x \$ 0.00
Independent claims	1-3 =		x \$ 0.00
MULTIPLE DEPENDENT CLAIM(s) (if applicable)			+ 280.00 \$ 280.00
TOTAL OF ABOVE CALCULATIONS =			\$ 1,320.00
<input type="checkbox"/> Applicant claims small entity status. See 37 CFR 1.27. The fees indicated above are reduced by 1/2.			\$ _____
SUBTOTAL =			\$ 1,320.00
Processing fee of \$ _____ for furnishing the English translation later than <input type="checkbox"/> 20 <input type="checkbox"/> 30 months from the earliest claimed priority date (37 CFR 1.492 (f)).			\$ _____
TOTAL NATIONAL FEE =			\$ 1,320.00
Fee for recording the enclosed assignment (37 CFR 1.21 (h)) Assignment must be accompanied by appropriate cover sheet (37 CFR 3.28, 3.31) (_____ per property).			+ \$ _____
TOTAL FEES ENCLOSED =			\$ 1,320.00
			Amount to be Refunded: \$ _____
			Charged: \$ _____
a. <input type="checkbox"/> A check in the amount of \$ _____ to cover the above fees is enclosed. b. <input checked="" type="checkbox"/> Please charge my Deposit Account No. 12-1095 in the amount of \$ 1,320.00 to cover the above fees. A duplicate copy of this sheet is enclosed. c. <input checked="" type="checkbox"/> The Commissioner is hereby authorized to charge any additional fees which may be required or credit any overpayment to my Deposit Account No. 12-1095. A duplicate copy of this sheet is enclosed.			
NOTE: Where an appropriate time limit under 37 CFR 1.494 or 1.495 has not been met, a petition to revive (37 CFR 1.137 (a) or (b)) must be filed and granted to restore the application to pending status.			
SEND ALL CORRESPONDENCE TO Arnold H. Krumholz LERNER, DAVID, LITTENBERG, KRUMHOLZ & MENTLIK, LLP 600 South Avenue West Westfield, New Jersey 07090 (908) 518-6304			
 SIGNATURE			Arnold H. Krumholz
NAME			
			25,428
REGISTRATION NUMBER			

i/pitv

Process for preventing scaling of heat transfer surfaces.**Background of the invention**

- In the kraft cooking process, cellulosic material, most conveniently in the form of chips, is treated at elevated temperatures, typically 160-180 °C, with alkaline cooking liquor containing sodium hydroxide and sodium sulfide. The fresh inorganic liquor is referred to as white liquor, and the spent liquor containing the dissolved wood material is referred to as black liquor.
- Since the emergence of the kraft cooking process until the present date, one of the most important objectives has been the attempt to reduce the energy consumption of the cooking process. Processes have therefore been developed for the purpose of, among other aspects, energy saving. In continuous processes, this may take place by heating the chip material with secondary steam obtained from flashing the hot black liquor. In batch cooking, the most useful technique is to recover the hot black liquor at the end of cooking and reuse its energy 1) as a direct heating medium to be pumped into the digester during a subsequent batch, and 2) to heat up white liquor by means of heat exchangers. Good examples of this development are batch processes described in, e.g. Fagerlund, U.S. Pat No 5,578,149 and Östman, U.S. Pat. 4,764,251. The displaced liquors of over 100°C are stored in one or several pressurized accumulators which further contain a continuous heat recovery system (see, e.g. U.S. Pat. No. 5,643,410). As a result, the energy efficiency of batch cooking has increased.

- Another important objective has been to improve the properties and quality of the pulp produced. In the liquor displacement batch method, avoiding digester discharge by means of hard hot blow techniques, has made this possible. Gentle digester discharge is typically accomplished by cooling the digester prior to discharge, relieving the overpressure in the digester and then pumping the cooked material from the digester (see, e.g., U.S. Pat. 4,814,042). Further development of liquor-displacement kraft batch cooking has also involved the combination of energy efficiency and efficient usage of residual and fresh cooking chemicals to facilitate delignification and high pulp strength (see, e.g., U.S. Pat. No. 5,183,535 and U.S. Pat. No. 5,643,410). This can be accomplished by arranging the

displacement at the end of the cook to first recover the "mother" black liquor, hot and rich in residual sulfur, into one accumulator and then to recover the portion of black liquor contaminated by wash filtrate and lower in solids and temperature in another accumulator. The accumulated black liquors are then reused in reverse order to impregnate and react with, respectively, the next batch of wood chips prior to finalization of the cook with white liquor. Thus it has become possible to start a kraft cook with a high charge of sulfur and a low charge of alkali and thus carry out important sulfur-lignin reactions in the early phase, especially in the hot black liquor treatment, facilitating subsequent delignification with fresh cooking liquor.

Thus, the above-mentioned development of the batch cooking technology has been characterized by improvements in terms of energy savings and properties of the delignified cellulosic material as e.g. strength and uniformity.

Important in the cooking process is also that the process has a good fit to surrounding processes as e.g. spent liquor evaporation and pulp washing. Through black liquor evaporation, incineration, melting of the smelt into a water solution and causticizing the resulting liquor, white liquor is regenerated from the chemicals contained in the black liquor. This is the basis for recovery of alkaline spent liquors.

Traditionally, both in the case of batch cooking processes and continuously operated cooking processes, the black liquor led to the evaporation originates from the main cooking stage at elevated temperature. In the search for improved energy efficiency and improved properties of the delignified cellulosic material, cooking methods have been developed wherein the black liquor fed to the evaporation plant is recycled black liquor originating from the early stages of the cooking sequence. Such processes have been disclosed in e.g., U.S. Pat. No. 5,643,410.

It has been observed that the properties of the black liquor originating from the early stages of the cook differ from those of black liquor from a traditional cook. Recycled black liquor originating from the early stages of the cooking sequence may complicate the evaporation of black liquor. A particular problem is fouling of the surfaces of heat exchangers in the

evaporation plant, leading to a decrease in heat transfer. Fouling may be so extensive that the heat transfer surfaces must be repeatedly cleaned, which requires special procedures, calls for an evaporation plant shutdown and may even limit production. The evaporator fouling problems with black liquors that originates from the early stages of cooking are typically related to calcium.

In the early stage of alkaline cooking, calcium-containing material dissolves into the black liquor from the lignocellulosic material. In a traditional cook, the cook proceeds by heating, the temperature increases and no essential liquor exchange occurs. Then, a major part of the dissolved calcium-containing material in the cooking liquor is broken down, calcium carbonate is formed and as a result, a major part of the calcium is resorbed onto the lignocellulosic material in the digester. Typically, following such a cooking process, evaporation of the black liquor can be carried out without problems caused by precipitation of calcium on the heat transfer surfaces, as the black liquor fed to evaporation originates from the cooking stage at elevated temperature.

The evaporation problems with black liquors originating from the early stages of cooking, are typically related to such calcium-containing material dissolved in the early stages of a cook. The dissolved calcium-containing material has not been degraded, and the amount of calcium bound to the dissolved material in the black liquor is high. In subsequent evaporation processes, the solids content of the black liquor rises and the evaporation temperature typically increases. Thus, the dissolved material is degraded further as evaporation proceeds and the solids content and temperature both rise. The calcium bound to the dissolved material in the black liquor is thereby set free. The liberated calcium reacts with the carbonate in the black liquor, forming calcium carbonate. A significant amount of crystallization occurs on the evaporation plant's heat transfer surfaces, whereby the plant's water evaporation capacity is severely limited. Crystallization may be so extensive, that the heat transfer surfaces must be repeatedly cleaned.

It has been shown that a liquor's potential for calcium carbonate precipitation is dependent on the temperature history of the liquor. Fredrick and Grace¹ (Southern Pulp and Paper Manufacturer 42 (1979) 8:16-23) have proposed that the amount of dissolved calcium is

increased because in the black liquor the calcium forms a complex with the lignin. This complex is unstable at higher temperatures. At higher temperatures, the complex breaks down, and if calcium ions are released close to a hot surface, the calcium ion reacts with carbonate ions present in the liquor and precipitate is formed on the surface.

5

Frederick and Grace have proposed that calcium precipitation can be decreased or avoided by heating the black liquor between evaporation stages to temperatures of 150-160°C and times of 10-20 minutes. However, the above mentioned method has not been extensively in use because it raises investment and operating costs.

10

Magnusson, Sjölander and Liden have suggested, that the existence of dissolved calcium in kraft black liquors can be explained by the high amounts of dissolved carbon present (Tappi 1998 International Chemical Recovery Conference Proceedings, Tampa, FL, USA, 1-4 June 1998, Vol. 1, p. 379-383). The introduction of dissolved forms of calcium into the black liquor will lead to an increased degree of supersaturation. Since this form of calcium is thermodynamically unstable with respect to calcium carbonate formation, heating of such liquors will, at some elevated temperature, cause rapid precipitation. They showed that calcium carbonate precipitation occurs in the temperature range of 110-145°C, and they also suggested that potential danger of scaling problems in evaporation plants could be avoided by heat treatment that triggers calcium carbonate precipitation during a process stage where it does not lead to harmful scaling.

15

20

25

Others have proposed direct heating of black liquors fed to evaporator at a temperature of 110-145°C and times of 1-20 minutes (patent application FI 980387). However, also this method increases investment costs.

Disclosure of the invention

30

In accordance with the present invention, these and other objects has now been accomplished by the discovery of a new method for preparing pulp. More particularly, the present invention relates to processes for preparing pulp in which cellulosic material is treated in one or several impregnation and pretreatment stages before delignifying the heated cellulosic material with fresh alkaline cooking liquor at elevated temperatures. The present in-

vention specifically relates to the liquor exchange in a cooking process that produces high quality pulp, is energy efficient and generates spent liquor, which liquor can be processed in the evaporation plant without forming scaling of low solubility.

- 5 For the purpose of this discussion, "calcium-containing spent liquor" refers to a process liquor containing calcium bound to dissolved organic material. The calcium content is calculated on the basis of calcium in the dry solids content of the liquor.

- 10 In accordance with the present invention, a process is provided for the preparation of pulp from lignin-containing cellulosic material using alkaline cooking, which process comprises the stages of a) charging lignocellulose-containing material to a digester, b) treating said lignocellulose-containing material initially with an impregnation liquor, and subsequently with hotter liquors, displacing calcium-containing spent liquor from the digester, c) heating and cooking said lignocellulose-containing material at cooking temperature and pressure to
15 produce cooked lignocellulose-containing material and cooking liquor, d) displacing said cooking liquor using at least part of said calcium-containing spent liquor, whereby said calcium-containing spent liquor is heated by the digester contents .

- 20 Preferably, stage b) includes the sequential introduction of impregnation liquor, hot black liquor and preheated white liquor in proper ratios. The last part of the final displacement stage in d) is preferably carried out using wash filtrate from the downstream process.

- In accordance with one embodiment of the process of the present invention, the temperature of the impregnation liquor is between about 20 and 100°C, the temperatures of the
25 hotter liquors are between about 120 and 180 °C and the temperature of displaced calcium-containing spent liquors are between about 20 and 160 °C.

- In accordance with one embodiment of the process of the present invention, the method includes monitoring the calcium content of spent liquor being displaced, in order to determine the proper cut points for isolating displaced calcium-containing spent liquor which
30 may cause calcium precipitation at higher temperatures and/or dry solid contents.

In accordance with another embodiment of the process of the present invention, the method includes monitoring the temperature of spent liquor being displaced, in order to determine the proper cut points for isolating displaced calcium-containing spent liquor.

- 5 In accordance with another embodiment of the process of the present invention, the temperature of displaced calcium-containing spent liquor is between about 20 and 100°C.

- In accordance with another embodiment of the present invention, a process using alkaline cooking is provided for the preparation of pulp from lignin-containing cellulosic material, which process comprises a) charging lignocellulose-containing material to a digester; b) treating said lignocellulose-containing material initially with an impregnation liquor and subsequently with hotter liquors, displacing from the digester during said subsequent treatments a first portion of calcium-containing spent liquor, and at least a second portion of calcium-containing spent liquor at lower calcium content compared to said first portion of calcium-containing spent liquor, c) heating and cooking said lignocellulose-containing material to produce cooked lignocellulose-containing material and cooking liquor, d) displacing said cooking liquor initially with said first portion of calcium-containing spent liquor, whereby said calcium-containing spent liquor is heated by the digester contents, and subsequently with wash filtrate so as to displace spent liquor and cool the digester content; and e) discharging the digester.
- 10
15
20

- In accordance with another embodiment of the present invention, several portions of calcium-containing spent liquor containing different amounts of calcium are used for displacement of cooking liquor, sequentially or in combination with wash filtrate.
- 25

- In accordance with the present invention, a process using alkaline cooking is provided for the preparation of pulp from lignin-containing cellulosic material, said process comprising the steps of a) charging lignocellulose-containing material to a digester; b) treating said lignocellulose-containing material initially with an impregnation liquor and subsequently with hotter liquors, thereby displacing from the digester a first portion of calcium-containing spent liquor and a second portion of calcium-containing spent liquor having a lower calcium content compared to said first portion of calcium-containing spent liquor; c)
- 30

heating and cooking said lignocellulose-containing material to produce cooked lignocellulose-containing material and cooking liquor; d) displacing said cooking liquor initially with a portion of said calcium-containing spent liquor, whereby said calcium-containing spent liquor is heated by the digester contents, and subsequently with wash filtrate so as to displace a first portion of cooking liquor, said first portion of cooking liquor having a temperature and dry solids content substantially corresponding to the temperature and dry solids content of the cooking liquor at the end of the cook; and to displace a second portion of cooking liquor, said second portion of cooking liquor having a temperature and dry solids content substantially lower than the temperature and dry solids content of the cooking liquor at the end of the cook; maintaining the first and second portion of cooking liquor separate from each other; combining the second portion of calcium-containing liquor and the second portion of cooking liquor; utilizing the first portion of cooking liquor for pretreating and heating the lignocellulosic material of a subsequent batch; and using the combined second portion of calcium-containing liquor and second portion of cooking liquor for supplying heat to a subsequent batch of lignocellulose-containing material.

In a preferred embodiment of the present invention, the method includes transferring the combined second portion of calcium-containing liquor and second portion of cooking liquor, after supplying heat to the subsequent batch of lignocellulose-containing material, to a liquor tank at atmospheric pressure. Preferably, the method includes separating and removing soap contained in said combined portions of liquor.

In a preferred embodiment of this method of the present invention, the method includes transferring the combined second portion of calcium-containing liquor and second portion of cooking liquor, after supplying heat to a subsequent batch of lignocellulose-containing material, to an evaporation plant for recovery of cooking chemicals.

In accordance with one embodiment of this method of the present invention, the method includes utilizing wash filtrate and a combined second portion of calcium-containing liquor and second portion of cooking liquor and for impregnating the lignocellulose-containing material in an impregnation step.

In a preferred embodiment, the wash filtrate comprises a filtrate from a subsequent wash plant for kraft pulp.

5 In accordance with another embodiment of this method of the present invention, the method includes utilizing the combined second portion of calcium-containing liquor and second portion of cooking liquor for preheating fresh alkaline cooking liquor supplied to the digester.

10 As calcium-containing spent liquor is fed into the digester following the cooking stage, when the cooking liquor at cooking temperature is displaced, the calcium-containing material extracted in the early stages of the process decomposes when the calcium-containing liquor enters the hot digester. Thus, the calcium bound to dissolved material is set free in the digester because of the high temperature. The liberated calcium reacts with carbonate ion, and calcium carbonate is formed in the digester where no scaling can be formed on
15 heat transfer surfaces. Thus, the formed calcium carbonate is retained in the pulp and is not extensively carried to the evaporating plant.

In general, the present invention provides a method for overcoming a drawback in prior art low energy kraft batch cooking processes. In the process according to the invention, kraft
20 pulp is prepared and spent liquor is generated which, when fed to an evaporation plant, can be processed without forming scaling of low solubility. An essential advantage is that the method does not require any essential investments and has no impact on sequence times and production capacity.

25

Detailed description

Figure 1 shows a block diagram of a liquor-displacement kraft batch process according to the present invention. The figure defines the required tanks, streams and the cooking sequence. Charging the digester with wood chips and evacuating the digester starts the kraft
30 cook. The chips can be packed with steam or be pre-steamed, before the digester is filled essentially with impregnation liquor A from the impregnation liquor tank 5, soaking and heating the chips.. Wood chip charging and impregnation liquor charging preferably over-

lap. An overflow, A1, to black liquor tank, point AB, is carried out in order to remove air and diluted first front of liquor. Preferably, the volume of A1 is kept low. After closing the flow, the digester is pressurized and impregnation is completed. During impregnation, a relatively low temperature is preferred, since a higher impregnation temperature will consume residual alkali too fast, resulting in higher rejects and non-uniform cooking. Preferably, the temperature of this impregnation step is below 100°C. In practice, temperatures of from about 20 °C to 100 °C can be utilized.

In the next stage, the wood chips are further treated with hotter liquors before actual cooking. The temperatures of the hotter liquors are between 120 °C to 180 °C. In figure 1, a method is described where hot black liquor B is pumped in from hot black liquor tank 1. Black liquor from tank 1 is at constant temperature, dry solids content and residual alkali, which makes it easy to maintain conformity from cook to cook. This is extremely important because the hot black liquor has a major chemical effect on the wood, and controls the selectivity and cooking kinetics in the main cooking phase with white liquor. The cooler and partly diluted impregnation liquor A2, displaced by hot black liquor, is essentially conducted to black liquor tank 4, point AB. In the spent liquor portions A1 and A2, the dissolved material typically has the highest content of calcium. The cooking sequence is continued by pumping in hot white liquor, point C, from the hot white liquor tank 3 and a smaller amount of hot black liquor, B, 1) simultaneously with the hot white liquor, in order to dilute the very high alkali concentration of fresh white liquor and 2) after white liquor charge, in order to flush lines into the digester. The liquor D2, displaced by hot liquor, preferably above about the atmospheric boiling point, is conducted to hot black liquor tank 2.

According to the prior art as disclosed in e.g. U.S. Pat. No. 5,643,410, the displaced liquors A1 and A2 are essentially conducted to the evaporation plant. This procedure will however transfer the calcium-containing dissolved material, which has not yet been degraded in the cooking process, to the evaporation plant. In the subsequent evaporation process, during which the solids content and temperature typically increase, the calcium bound to the dissolved material in the black liquor is set free. As a consequence, calcium carbonate scaling

occurs on heat transfer surfaces of evaporation units, whereby the plant's evaporation capacity is severely impaired.

According to the present invention, tank 4 is for storing the spent cooking liquor portions containing released calcium-containing material from the early impregnation and pretreatment stages of the cooking sequence, which liquors tend to form calcium precipitates at higher temperatures and dry solid contents. Preferably, the portion at the highest concentration of calcium is stored in tank 4. Most preferably, tank 4 is an atmospheric tank. The exact volume to be recovered to tank 4 is most suitably controlled by monitoring the calcium content, dry solids concentration and temperature of the displaced liquor exiting from the digester. After detecting a clear drop in calcium content bound to the dissolved material, the displaced liquor is switched to enter black liquor tank 2. The switch preferably occurs before the atmospheric boiling point of the displaced liquor is exceeded.

Further division of the calcium-containing displaced liquor is also possible, depending on available tank capacity. Preferably, the fractions are then used for final displacement in the same order, as described later.

After the filling procedure described above, the digester temperature is close to the final cooking temperature. The final cooking temperature can be between about 140 °C to 180 °C depending on the wood raw material. The final heating-up is carried out using direct or indirect steam heating and digester re-circulation. During cooking, additional fresh cooking liquor, C, from tank 3 can be added to even out the alkali profile. Spent liquor, B2, is then removed from the digester to tank 1 or tank 2.

After the desired cooking time, when delignification has proceeded to the desired reaction degree, the now spent cooking liquor is ready to be displaced with cool liquor, which serves to stop the cooking reactions and to cool and wash the digester content. According to the invention, the cooking liquor is at least partly displaced with a liquor portion E from the black liquor tank 4. Preferably, portion E is initially introduced followed by washing filtrate, point F, from the wash filtrate tank 6. In this manner, the calcium-containing liquors generated in the early stages of the process, i.e. portion E from tank 4, which contains

the highest amount of calcium bound to dissolved material, and is most liable to cause scaling precipitates in evaporation according to prior art methods, is introduced in the digester as the digester contents are at a high initial temperature. Consequently, the dissolved material decomposes and calcium carbonate is formed mainly within the bulk of the liquor, and remains in the liquor as calcium carbonate crystals or is absorbed into the pulp. Thereby, the spent liquor conducted to the evaporation plant from the cooking process is essentially free of problematic calcium bound to dissolved material.

If further subdivision of the calcium-containing liquors displaced from the digester has taken place, the fractions may be used in the same order for the final displacement, the latter fractions of successively lower calcium content experiencing successively lower temperatures.

No prior art technology using the impregnation and pretreatment procedures described above is able to produce spent liquor for the evaporation plant without calcium carbonate-related problems. The black liquor tank 4 now has a new role in the cooking process. It first collects the most problematic calcium-containing spent liquors and then transfers these liquors to the terminal displacement of the hot cooking liquor, preferably to the initial stages of the terminal displacement.

The invention will not essentially affect the washing efficiency of the terminal displacement, since the portion E contains chip water and some diluted spent liquor. Thus, the dissolved solids content of portion E is in the same order as that of the washing filtrate F. The cooling efficiency of the terminal displacement will be improved since the portion E has a low temperature, preferably 20 to 100 °C.

In the terminal displacement, the first portion B1 of exiting hot black liquor corresponds to the total of the volumes B required in the filling stages. Preferably, the terminal displacement is carried out to produce a displaced portion of spent cooking liquor having a temperature and dry solids content substantially corresponding to the predetermined temperature and dry solids content of the cooking liquor. The second portion D1 of displaced spent cooking liquor, which is diluted by the displacement filtrate but is still above its atmospheric boiling point, is conducted to the hot black liquor tank 2. After completed final displacement

placement, the digester contents are discharged for further processing of the pulp. The above cooking sequence may then be repeated.

The equipment for the cooking process includes the tank farm, where fresh liquors and spent liquors are stored and heat is recovered. The impregnation liquor tank is provided with wash liquor from the washing plant. The hot black liquor tank 2 provides spent black liquor to the evaporation plant through fiber separation, and also partly provides cooled impregnation black liquor to the impregnation liquor tank, transferring heat to white liquor and water by means of heat exchange. The spent liquor transferred to the evaporation plant, according to the invention, thus originates from conditions at higher temperatures compared to prior art technology such as that described in U.S. Pat. No. 5,643,410. According to the invention, tank 2 also has a function of decomposing calcium-containing dissolved material, as tank 2 provides residence times of 10 to 60 minutes and temperatures from about 100 to 135 °C. The temperature of tank 2 depends, among other factors, on the arrangement of terminal displacement, temperature of charged lignocellulosic material, and the switching point from collecting to tank 4 to tank 2 in the pretreatment step. An additional advantage of the invention compared to prior art such as the process of U.S. Pat. No. 5,643,410, is that the temperature of the spent liquor conducted to the evaporation plant can be controlled. This is especially important when outside temperatures are low, e.g. during winter when temperatures in the lignocellulosic material charged to the cooking system is below the freezing point of water. In prior art processes, the temperature of the evaporation liquor has dropped in winter conditions, causing control difficulties and increased heating requirements in the evaporation plant. Another advantage of the invention is that the fiber separation of the spent liquor conducted the evaporation plant can occur at higher temperatures, facilitating soap solubility and reducing carryover of soap to the separated fiber fraction. The soap is preferably conducted with the spent liquor to the evaporation plant, where soap can be further removed.

Tank 4, 5 and 6 are furnished with soap separation equipment according to prior art soap separation technology. Practical experience on mill-scale has proven that soap removal in these locations of the black liquor transfer sequence is of major importance especially

when processing soap-containing softwood raw materials. It is of great importance to use only low-in-soap black liquor for impregnation or other digester filling purposes.

5 Although the invention herein has been described with reference to particular embodiments, it is to be understood that these embodiments are merely illustrative of the principles and applications of the present invention. It is therefore to be understood that numerous modifications may be made to the illustrative embodiments and that other arrangements may be devised without departing from the spirit and scope of the present invention as defined by the appended claims.

We claim:

1. A method for producing chemical pulp by means of alkaline cooking, comprising the steps of
 - 5 a) charging lignocellulose-containing material to a digester;
 - b) treating said lignocellulose-containing material initially with an impregnation liquor and subsequently with hotter liquors, displacing calcium-containing spent liquor from the digester during said treatment;
 - c) heating and cooking said lignocellulose-containing material at cooking temperature and
10 pressure so as to produce cooked lignocellulose-containing material and cooking liquor;
 - d) displacing said cooking liquor from the digester;**characterized** in that said cooking liquor is displaced using at least part of the displaced calcium-containing spent liquor from stage b).
- 15 2. The method according to claim 1, **characterized** by the calcium-containing spent liquor in step b) being displaced and collected into at least two separate portions, the latter portion(s) having lower calcium content, on a dry solids basis, than the first portion.
3. The method according to claim 2, **characterized** by a latter portion of calcium-
20 containing spent liquor being combined with a portion of displaced cooking liquor and the combined liquors being used for supplying heat to a subsequent batch.
5. The method according to claim 2, **characterized** by the first portion of displaced calcium-containing spent liquor being used for displacement of cooking liquor after the
25 cooking stage.
6. The method according to claim 5, **characterized** by said first portion being the first portion of liquor introduced into the digester for displacement.
7. The method according to claim 2, **characterized** by at least one of the latter portions of calcium-containing spent liquor being used for displacement of cooking liquor.

8. The method according to any claim 1-7, **characterized** by the calcium content, on a dry solids basis, of the calcium-containing spent liquor being monitored during its displacement.
- 5
9. The method according to any claim 1-7, **characterized** by the temperature of the calcium-containing spent liquor being monitored during its displacement.

(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(19) World Intellectual Property Organization
International Bureau(43) International Publication Date
12 April 2001 (12.04.2001)

PCT

(10) International Publication Number
WO 01/25531 A1

- (51) International Patent Classification⁷: D21C 3/22, 11/10
- (21) International Application Number: PCT/FI00/00855
- (22) International Filing Date: 4 October 2000 (04.10.2000)
- (25) Filing Language: English
- (26) Publication Language: English
- (30) Priority Data:
19992132 4 October 1999 (04.10.1999) FI
- (71) Applicant (for all designated States except US): VALMET
CHEMICAL PULPING OY [FI/FI]; P.O. Box 34, FIN-
28101 Pori (FI).
- (72) Inventors; and
(73) Inventors/Applicants (for US only): PAAKKI, Asko
[FI/FI]; Ahonkuja 7, FIN-53300 Lappeenranta (FI), UUSI-
TALO, Päivi [FI/FI]; Hepokuja 6, FIN-29600 Noormarkku
(FI), SVEDMAN, Mikael [FI/FI]; Itsenäisyydenkatu 66
as 4, FIN-28100 Pori (FI).
- (74) Agent: OY JALO ANT-WUORINEN AB; Iso
Roobertinkatu 4-6 A, FIN-00120 Helsinki (FI).
- (81) Designated States (national): AE, AG, AL, AM, AT, AT
(utility model), AU, AZ, BA, BB, BG, BR, BY, BZ, CA,
CH, CN, CR, CU, CZ, CZ (utility model), DE, DE (utility
model), DK, DK (utility model), DM, DZ, EE, EE (utility
model), ES, FI, FI (utility model), GB, GD, GE, GH, GM,
HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KR (utility
model), KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG,
MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD,
SE, SG, SI, SK, SK (utility model), SL, TJ, TM, TR, TT,
TZ, UA, UG, US, UZ, VN, YU, ZA, ZW.
- (84) Designated States (regional): ARIPO patent (GH, GM,
KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW), Eurasian
patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European
patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE,
IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG,
CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).

Published:

- With international search report.
- Before the expiration of the time limit for amending the claims and to be republished in the event of receipt of amendments.

For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

(54) Title: PROCESS FOR PREVENTING SCALING OF HEAT TRANSFER SURFACES

(57) Abstract: In a batch process for kraft pulp production, fouling of heat transfer surfaces is reduced. The first impregnation liquor withdrawn from the digester is rich in dissolved calcium-containing material. This material, when heated, releases calcium which forms calcium carbonate. If this heating takes place at a heat exchanger surface, severe scaling may result. According to the invention, the calcium-rich liquor is introduced into the digester as an early displacement liquor following a finished cook. As the digester temperature is high at this process stage, the calcium-containing dissolved material breaks down and calcium carbonate precipitation takes place in the pulp in the digester, where scaling of heat transfer surfaces cannot occur.

1/1

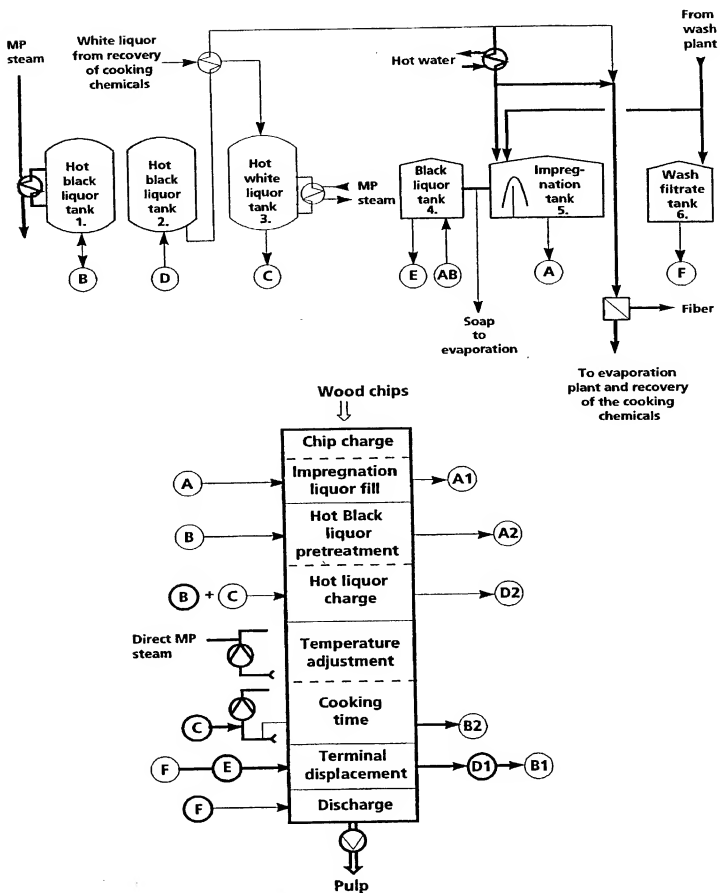


Figure 1

Please type a plus sign (+) inside this box



PTO/SB/01 (03-01)

Approved for use through 10/31/2002 OMB 0651-0032

U.S. Patent and Trademark Office; U.S. DEPARTMENT OF COMMERCE

Under the Paperwork Reduction Act of 1995, no persons are required to respond to a collection of information unless it contains a valid OMB control number

DECLARATION FOR UTILITY OR DESIGN PATENT APPLICATION (37 CFR 1.63)

☐ Declaration
Submitted
with Initial
Filing

OR

☐ Declaration
Submitted after Initial
Filing (surcharge
(37 CFR 1.16 (e))
required)

Attorney Docket Number OYJALO-011

First Named Inventor Aiko PAAKKI

COMPLETE IF KNOWN

Application Number

Filing Date

Group Art Unit

Examiner Name

As a below named inventor, I hereby declare that:

My residence, mailing address, and citizenship are as stated below next to my name.

I believe I am the original and first inventor of the subject matter which is claimed and for which a patent is sought on the invention entitled:

PROCESS FOR PREVENTING SCALING OF HEAT TRANSFER SURFACES

(Title of the Invention)

the specification of which

☐ is attached hereto

OR

☒ was filed on (MM/DD/YYYY) 10/04/00 as United States Application Number or PCT International

Application No. PCT/FI00/00859 and was amended on (MM/DD/YYYY) (if applicable).

I hereby state that I have reviewed and understand the contents of the above identified specification, including the claims, as amended by any amendment specifically referred to above.

I acknowledge the duty to disclose information which is material to patentability as defined in 37 CFR 1.56, including for continuation-in-part applications, material information which became available between the filing date of the prior application and the national or PCT international filing date of the continuation-in-part application.

I hereby claim foreign priority benefits under 35 U.S.C. 119(a)-(d) or (f), or 365(b) of any foreign application(s) for patent, inventor's or plant breeder's rights certificate(s), or 365 (a) of any PCT international application which designated at least one country other than the United States of America, listed below and have also identified below, by checking the box, any foreign application for patent, inventor's or plant breeder's rights certificate(s), or of any PCT international application having a filing date before that of the application on which priority is claimed.

Prior Foreign Application Number(s)	Country	Foreign Filing Date (MM/DD/YYYY)	Priority Not Claimed	Certified Copy Attached?	
				YES	NO
1992132 ✓	Finland ✓	10/04/99 ✓	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
			<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
			<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
			<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

☐ Additional foreign application numbers are listed on a supplemental priority data sheet PTO/SB/02B attached hereto:

Please type a plus sign (+) inside this box



PTO/SB/01 (03-01)

Approved for use through 10/31/2002. OMB 0651-0032

U.S. Patent and Trademark Office, U.S. DEPARTMENT OF COMMERCE

Under the Paperwork Reduction Act of 1995, no persons are required to respond to a collection of information unless it contains a valid OMB control number

DECLARATION — Utility or Design Patent Application

POWER OF ATTORNEY As a named inventor, I hereby appoint the following registered practitioner(s) to prosecute this application and to transact all business in the Patent and Trademark Office connected therewith. Customer Number 000530

Direct all correspondence to

☐ Customer Number
or Bar Code Label
OR ☐ Correspondence address below

00530
PATENT-TRADEMARK OFFICE

Name

Address

City

State

ZIP

Country

Telephone

Fax

I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under 18 U.S.C. 1001 and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

NAME OF SOLE OR FIRST INVENTOR:

☐ A petition has been filed for this unsigned inventor

Given Name
(first and middle (if any)) Asko

Family Name
or Surname Paakki

Inventor's
Signature *Asko Paakki*

Date March 11, 2002

Residence: City Lappeenranta

State FINCountry FINLANDCitizenship Finnish ✓

Mailing
Address:

Ahonkuja 7, 00City Lappeenranta

State

ZIP 53300Country FINLAND

NAME OF SECOND INVENTOR:

☐ A petition has been filed for this unsigned inventor

Given Name
(first and middle (if any)) Päivi

Family Name
or Surname Uusitalo

Inventor's
Signature *Päivi Uusitalo*

Date March 11, 2002

Residence: City Noormarkku

State FINCountry FINLANDCitizenship Finnish ✓

Mailing
Address:

Hepokuja 6City Noormarkku

State

ZIP 29600Country FINLAND☒

Additional inventors are being named on the

1

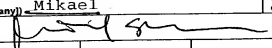
supplemental Additional Inventor(s) sheet(s) PTO/SB/02A attached hereto

Please type a plus sign (+) inside this box ☐

PTO/SB/02A (11-00)
 Approved for use through 10/31/2002. OMB 0651-0032
 U.S. Patent and Trademark Office; U.S. DEPARTMENT OF COMMERCE
 Under the Paperwork Reduction Act of 1995, no persons are required to respond to a collection of information unless it contains a valid OMB control number.

DECLARATION

 ADDITIONAL INVENTOR(S)
 Supplemental Sheet
 Page 1 of 1

Name of Additional Joint Inventor, if any:		<input type="checkbox"/> A petition has been filed for this unsigned inventor	
Given Name (first and middle (if any)) <u>Mikael</u>		Family Name or Surname <u>Svedman</u>	
Inventor's Signature 		Date <u>March 11, 2002</u>	
Residence: City <u>Pori</u> FD State	Country <u>FINLAND</u>	Citizenship <u>Finnish</u>	
Mailing Address:	<u>Itsenäisyydenkatu 66</u>		
City <u>Pori</u>	State	ZIP <u>28100</u>	Country <u>FINLAND</u>
Name of Additional Joint Inventor, if any:		<input type="checkbox"/> A petition has been filed for this unsigned inventor	
Given Name (first and middle (if any))		Family Name or Surname	
Inventor's Signature		Date	
Residence: City	State	Country	Citizenship
Mailing Address:			
City	State	ZIP	Country
Name of Additional Joint Inventor, if any:		<input type="checkbox"/> A petition has been filed for this unsigned inventor	
Given Name (first and middle (if any))		Family Name or Surname	
Inventor's Signature		Date	
Residence: City	State	Country	Citizenship
Mailing Address:			
City	State	ZIP	Country
Name of Additional Joint Inventor, if any:		<input type="checkbox"/> A petition has been filed for this unsigned inventor	
Given Name (first and middle (if any))		Family Name or Surname	
Inventor's Signature		Date	
Residence: City	State	Country	Citizenship
Mailing Address:			
City	State	ZIP	Country